AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

LISTING OF CLAIMS:

1-26. (canceled)

27. (previously presented) A method for the treatment of Alzheimer's disease (AD) in a subject having or suspected of having AD, comprising administering to said subject a therapeutically effective amount of an antibody wherein said antibody is raised against a protofibril comprising an Aβ-Arc peptide selected from the group consisting of Aβ39-Arc (Amino Acids 1-39 of SEQ ID NO:1), Aβ40-Arc (Amino Acids 1-40 of SEQ ID NO: 1), and Aβ42-Arc (SEQ ID NO: 1), wherein said antibodies bind to arctic and wild-type Aβ peptides in protofibril conformation.

28-43. (canceled)

44. (previously presented) A method for the treatment of Alzheimer's disease (AD) in a subject having or suspected of having AD, comprising administering to said subject a therapeutically effective amount of an antibody, wherein said antibody is raised against a protofibril comprising an Aβ-Arc peptide selected from the group consisting of Aβ39-Arc (Amino Acids 1-39 of SEQ ID NO:1), Aβ40-Arc (Amino Acids 1-40 of SEQ ID NO:1), Aβ41-Arc (Amino Acids 1-41 of SEQ ID NO:1), Aβ42-Arc (SEQ

ID NO:1), and combinations thereof, wherein said antibodies bind to arctic and wild-type $A\beta$ peptides in protofibril conformation.

45. (previously presented) The method according to claim 44, wherein said protofibril further comprises an $A\beta$ peptide having a mutation selected from the group consisting of the Dutch, Flemish, Italian, Iowa mutations, and combinations thereof.

46-48. (canceled)

- 49. (previously presented) The method according to claim
 44, wherein said antibody is monoclonal.
- 50. (previously presented) The method according to claim 44, wherein said antibody is human or humanized.
- 51. (currently amended) A method for the treatment of Alzheimer's disease (AD) in a subject having or suspected of having AD, comprising administering to said subject a therapeutically effective amount of an antibody, wherein said antibody is raised against a composition comprising a protofibril comprising an Aβ-Arc peptide selected from the group consisting of Aβ39-Arc (Amino Acids 1-39 of SEQ ID NO:1), Aβ40-Arc (Amino Acids 1-40 of SEQ ID NO:1), Aβ41-Arc (Amino Acids 1-41 of SEQ ID NO:1), Aβ42-Arc (SEQ ID NO:1), and combinations thereof, wherein said antibodies bind to arctic and wild-type Aβ peptides in protofibril conformation.

52. (previously presented) The method according to claim 44, wherein said protofibril further comprises an $A\beta$ peptide having a Dutch mutation.

53-56. (canceled)

- 57. (previously presented) The method according to claim 44, wherein said A β -Arc peptide is A β 39-Arc (Amino Acids 1-39 of SEQ ID NO:1).
- $_{58.}$ (previously presented) The method according to claim 44, wherein said Aß-Arc peptide is Aß40-Arc (Amino Acids 1-40 of SEQ ID NO:1).
- 59. (previously presented) The method according to claim 44, wherein said A β -Arc peptide is A β 41-Arc (Amino Acids 1-41 of SEQ ID NO:1).
- 60. (previously presented) The method according to claim 44, wherein said A β -Arc peptide is A β 42-Arc (SEQ ID NO:1).
- 61. (new) The method according to claim 51, wherein said antibody is monoclonal.
- 62. (new) The method according to claim 51, wherein said antibody is human or humanized.
- 63. (new) The method according to claim 51, wherein said protofibril further comprises an $A\beta$ peptide with a mutation selected from the group consisting of the Dutch, Flemish, Italian and Iowa mutations.

- 64. (new) The method according to claim 51, wherein said A β -Arc peptide is A β 39-Arc (Amino Acids 1-39 of SEQ ID NO:1).
- 65. (new) The method according to claim 51, wherein said A β -Arc peptide is A β 40-Arc (Amino Acids 1-40 of SEQ ID NO:1).
- 66. (new) The method according to claim 51, wherein said A β -Arc peptide is A β 42-Arc (SEQ ID NO:1).
- 67. (new) The method according to claim 51, wherein said A β -Arc peptide is A β 41-Arc (Amino Acids 1-41 of SEQ ID NO:1).